

10724683 11/27/05

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID:SSSPTA1626KAS

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

\* \* \* \* \* Welcome to STN International \* \* \* \* \*

NEWS 1 Web Page URLs for STN Seminar Schedule - N. America  
NEWS 2 "Ask CAS" for self-help around the clock  
NEWS 3 SEP 09 ACD predicted properties enhanced in REGISTRY/ZREGISTRY  
NEWS 4 OCT 03 MATHDI removed from STN  
NEWS 5 OCT 04 CA/Caplus-Canadian Intellectual Property Office (CIPO) added  
to core patent offices  
NEWS 6 OCT 13 New CAS Information Use Policies Effective October 17, 2005  
NEWS 7 OCT 17 STN(R) AnaVist(TM), Version 1.01, allows the export/download  
of Caplus documents for use in third-party analysis and  
visualization tools  
NEWS 8 OCT 27 Free KWIC format extended in full-text databases  
NEWS 9 OCT 27 DIOGENES content streamlined  
NEWS 10 OCT 27 EPFULL enhanced with additional content  
NEWS 11 NOV 14 CA/Caplus - Expanded coverage of German academic research  
  
NEWS EXPRESS NOVEMBER 18 CURRENT VERSION FOR WINDOWS IS V8.01,  
CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),  
AND CURRENT DISCOVER FILE IS DATED 13 JUNE 2005.  
V8.0 USERS CAN OBTAIN THE UPGRADE TO V8.01 AT  
<http://download.cas.org/express/v8.0-Discover/>  
  
NEWS HOURS STN Operating Hours Plus Help Desk Availability  
NEWS INTER General Internet Information  
NEWS LOGIN Welcome Banner and News Items  
NEWS PHONE Direct Dial and Telecommunication Network Access to STN  
NEWS WWW CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that  
specific topic.

All use of STN is subject to the provisions of the STN Customer  
agreement. Please note that this agreement limits use to scientific  
research. Use for software development or design or implementation  
of commercial gateways or other similar uses is prohibited and may  
result in loss of user privileges and other penalties.

\* \* \* \* \* STN Columbus \* \* \* \* \*

FILE 'HOME' ENTERED AT 14:06:24 ON 27 NOV 2005

=> file reg

COST IN U.S. DOLLARS

SINCE FILE  
ENTRY

TOTAL  
SESSION

FULL ESTIMATED COST

0.21

0.21

FILE 'REGISTRY' ENTERED AT 14:06:39 ON 27 NOV 2005  
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.  
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.  
COPYRIGHT (C) 2005 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file  
provided by InfoChem.

STRUCTURE FILE UPDATES: 25 NOV 2005 HIGHEST RN 868737-05-7  
DICTIONARY FILE UPDATES: 25 NOV 2005 HIGHEST RN 868737-05-7

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH JULY 14, 2005

Please note that search-term pricing does apply when  
conducting SmartSELECT searches.

\*\*\*\*\*  
\*  
\* The CA roles and document type information have been removed from \*  
\* the IDE default display format and the ED field has been added, \*  
\* effective March 20, 2005. A new display format, IDERL, is now \*  
\* available and contains the CA role and document type information. \*  
\*  
\*\*\*\*\*

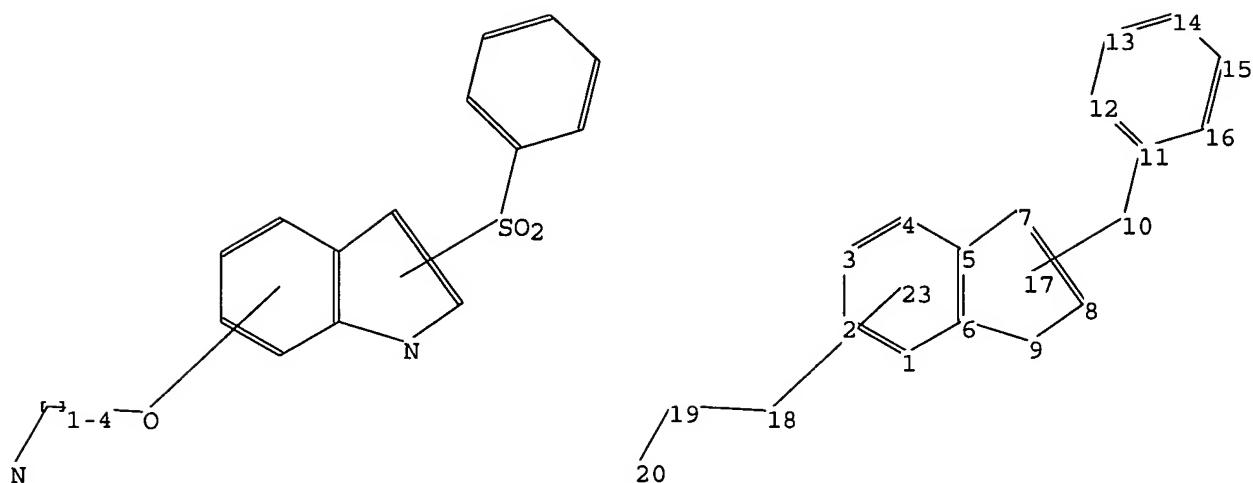
Structure search iteration limits have been increased. See HELP SLIMITS  
for details.

REGISTRY includes numerically searchable data for experimental and  
predicted properties as well as tags indicating availability of  
experimental property data in the original document. For information  
on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

=>

Uploading C:\Program Files\Stnexp\Queries\10724683.str



chain nodes :

10 18 19

ring nodes :

1 2 3 4 5 6 7 8 9 11 12 13 14 15 16

ring/chain nodes :

20

chain bonds :

10-11 18-19 19-20

ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-9 7-8 8-9 11-12 11-16 12-13 13-14 14-15  
15-16

exact/norm bonds :

5-7 6-9 7-8 8-9 18-19 19-20

exact bonds :

10-11

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6 11-12 11-16 12-13 13-14 14-15 15-16

isolated ring systems :

containing 11 :

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS

11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:CLASS 18:CLASS 19:CLASS

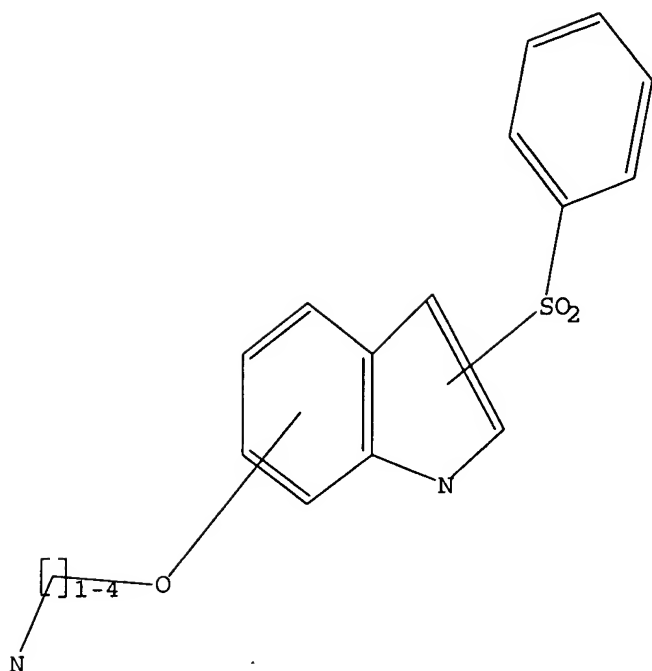
20:CLASS 23:CLASS

L1 STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s l1

SAMPLE SEARCH INITIATED 14:07:08 FILE 'REGISTRY'  
SAMPLE SCREEN SEARCH COMPLETED - 4610 TO ITERATE

43.4% PROCESSED 2000 ITERATIONS 2 ANSWERS  
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)  
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*  
BATCH \*\*COMPLETE\*\*  
PROJECTED ITERATIONS: 88129 TO 96271  
PROJECTED ANSWERS: 2 TO 220

L2 2 SEA SSS SAM L1

=> s l1 full

FULL SEARCH INITIATED 14:07:14 FILE 'REGISTRY'  
FULL SCREEN SEARCH COMPLETED - 93691 TO ITERATE

100.0% PROCESSED 93691 ITERATIONS 48 ANSWERS  
SEARCH TIME: 00.00.01

L3 48 SEA SSS FUL L1

=> file caplus

COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE	TOTAL
ENTRY	SESSION
161.33	161.54

10724683 11/27/05

FILE 'CAPLUS' ENTERED AT 14:07:21 ON 27 NOV 2005  
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.  
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.  
COPYRIGHT (C) 2005 AMERICAN CHEMICAL SOCIETY (ACS)

Copyright of the articles to which records in this database refer is held by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications. The CA Lexicon is the copyrighted intellectual property of the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

FILE COVERS 1907 - 27 Nov 2005 VOL 143 ISS 23  
FILE LAST UPDATED: 25 Nov 2005 (20051125/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/infopolicy.html>

=> s 13

L4                    2 L3

=> d ibib abs hitstr tot

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN  
 ACCESSION NUMBER: 2004:490721 CAPLUS  
 DOCUMENT NUMBER: 141:54192  
 TITLE: Preparation of aminoalkoxyindoles as 5-HT<sub>6</sub>-receptor ligands, in particular selective 5-HT<sub>6</sub> antagonists, for treating CNS disorders  
 INVENTOR(S): Zhao, Shu-Hai  
 PATENT ASSIGNEE(S): F. Hoffmann-La Roche A.-G., Switz.  
 SOURCE: PCT Int. Appl., 79 pp.  
 CODEN: PIXX22  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

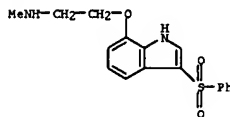
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004050085	A1	20040617	WO 2003-EP13372	20031127
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GR, GU, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, T, TH, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2508315	A1	20040617	CA 2003-2508315	20031127
EP 1569638	A1	20050907	EP 2003-782240	20031127
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
US 2004132799	A1	20040708	US 2003-724683	20031201
PRIORITY APPLN. INFO.:			US 2002-430506P	P 20021203
			WO 2003-EP13372	W 20031127
OTHER SOURCE(S):		MARPAT 141:54192		
GI				

\* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

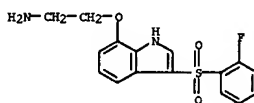
AB Title compds. I [wherein m = 0-3; n = 0-2; each R<sup>1</sup> = independently H, halo, halo/heteroalkyl, OH and derivs., NO<sub>2</sub>, CN, NH<sub>2</sub> and derivs., CONH<sub>2</sub> and derivs., SO<sub>2</sub>NH<sub>2</sub> and derivs., etc.; R<sup>2</sup> = (un)substituted heteroaryl; R<sup>3</sup> = H, alkyl; X = (CR<sup>4</sup>SR<sup>5</sup>)p; p = 2-3; R<sup>5</sup>, R<sup>6</sup>, R<sup>8</sup> = independently H, alkyl, or one of R<sup>5</sup> and R<sup>6</sup> together with one of R<sup>7</sup> and R<sup>8</sup> and the atoms therebetween = 4-6 membered heterocycle, etc.; and their pharmaceutically acceptable salts and prodrugs] were prepared as 5-HT<sub>6</sub>-receptor ligands, in particular selective 5-HT<sub>6</sub> antagonists, for treating CNS disorders. For example, I-HCl was prepared in 4 steps from 3-methyl-2-nitrophenol and 1-(2-chloroethyl)pyrrolidine hydrochloride by alkylation, cyclization, sulfonylation of the indole and oxidation with m-CPBA. Selected I displayed binding affinity values (pK<sub>i</sub>) for 5-HT<sub>6</sub> receptors in the range of 7.7 to 8.4 in an in vitro radioligand binding study. I were tested and found to

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)  
 N-[2-[(3-benzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]dimethylamine monohydrochloride 706783-75-7P, N-[2-[(3-(2-methoxybenzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]dimethylamine monohydrochloride 706783-76-8P, N-[2-[(3-(2-fluorobenzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]dimethylamine monohydrochloride 706783-77-9P, N-[2-[(3-(2-fluorobenzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]methylamine monohydrochloride 706783-78-0P, N-[2-[(3-benzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]methylamine monohydrochloride 706783-79-1P, N-[2-[(3-benzenesulfonyl)-1-methyl-1H-indol-7-yl]oxy]ethyl]methylamine monohydrochloride 706783-82-6P, N-[2-[(2-benzenesulfonyl)-1H-indol-4-yl]oxy]ethyl]methylamine monohydrochloride 706783-83-7P, N-[2-[(2-benzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]methylamine monohydrochloride 706783-84-8P, 3-(2,5-dichlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-85-9P, 2-benzenesulfonyl-4-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-87-1P, [2-[(3-(2-fluorobenzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]methylamine monohydrochloride 706784-06-7P, N-[2-[(3-(2-fluorobenzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]dimethylamine 706784-07-8P, 3-(2,5-dichlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-23-0P, 2-benzenesulfonyl-4-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-35-2P, 3-benzenesulfonyl-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-36-3P, 3-(3-chlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-37-4P, 3-(4-chlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-38-5P, 3-(2,3-dichlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-39-6P, 3-(2-chlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-40-9P, 3-(3,4-dichlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-41-0P, 3-(2-fluorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-42-1P, 3-(3-fluorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-43-2P, 3-(3-methoxybenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-44-3P, 3-(2-methoxybenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole 706784-45-4P, N-[2-[(3-benzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]dimethylamine 706784-46-5P, N-[2-[(3-(2-methoxybenzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]dimethylamine 706784-47-6P, N-[2-[(3-(2-fluorobenzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]methylamine 706784-48-7P, N-[2-[(3-benzenesulfonyl)-1-methyl-1H-indol-7-yl]oxy]ethyl]methylamine 706784-51-2P, N-[2-[(2-benzenesulfonyl)-1H-indol-4-yl]oxy]ethyl]methylamine 706784-52-3P, N-[2-[(2-benzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]methylamine  
 RI: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
 (5-HT<sub>6</sub> antagonist; prepn. of aminoalkoxyindoles as 5-HT<sub>6</sub>-receptor ligands, in particular selective 5-HT<sub>6</sub> antagonists, for treating CNS disorders)  
 RN 706783-64-4 CAPLUS  
 CN 1H-indole, 3-(phenylsulfonyl)-7-[2-(1-pyrrolidinyl)ethoxy]-, monohydrochloride (9CI) (CA INDEX NAME)

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)  
 be selective 5-HT<sub>6</sub> antagonists.  
 IT 706784-16-9P, N-[2-[(3-benzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]methylamine 706784-30-7P, [2-[(3-(2-fluorobenzenesulfonyl)-1H-indol-7-yl]oxy]ethyl]amine  
 RI: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)  
 (5-HT<sub>6</sub> antagonist; preparation of aminoalkoxyindoles as 5-HT<sub>6</sub>-receptor ligands, in particular selective 5-HT<sub>6</sub> antagonists, for treating CNS disorders)  
 RN 706784-16-9 CAPLUS  
 CN Ethanamine, N-methyl-2-[(3-(phenylsulfonyl)-1H-indol-7-yl]oxy]- (9CI) (CA INDEX NAME)

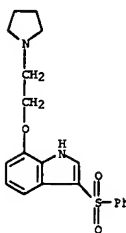


RN 706784-30-7 CAPLUS  
 CN Ethanamine, 2-[(3-[(2-fluorophenyl)sulfonyl]-1H-indol-7-yl]oxy]- (9CI) (CA INDEX NAME)



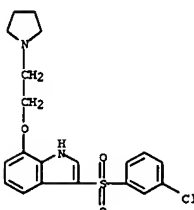
IT 706783-64-4P, 3-benzenesulfonyl-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-65-5P, 3-(3-chlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-66-6P, 3-(4-chlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-67-7P, 3-(2,3-dichlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-68-8P, 3-(2-chlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-69-9P, 3-(3,4-dichlorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-70-2P, 3-(2-fluorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-71-3P, 3-(3-fluorobenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-72-4P, 3-(3-methoxybenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-73-5P, 3-(2-methoxybenzenesulfonyl)-7-[2-(pyrrolidin-1-yl)ethoxy]-1H-indole monohydrochloride 706783-74-6P,

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)



● HCl

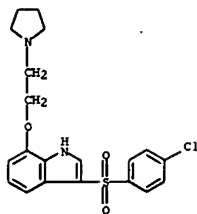
RN 706783-65-5 CAPLUS  
 CN 1H-indole, 3-[(3-chlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-, monohydrochloride (9CI) (CA INDEX NAME)



● HCl

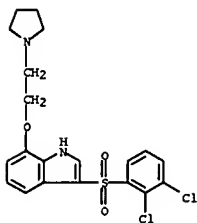
RN 706783-66-6 CAPLUS  
 CN 1H-indole, 3-[(4-chlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-, monohydrochloride (9CI) (CA INDEX NAME)

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)



● HCl

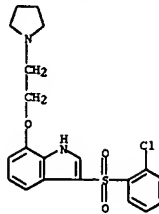
RN 706783-67-7 CAPLUS  
 CN 1H-Indole, 3-[(2,3-dichlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-,  
 monohydrochloride (9CI) (CA INDEX NAME)



● HCl

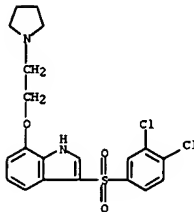
RN 706783-68-8 CAPLUS  
 CN 1H-Indole, 3-[(2-chlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-,  
 monohydrochloride (9CI) (CA INDEX NAME)

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)



● HCl

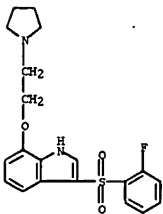
RN 706783-69-9 CAPLUS  
 CN 1H-Indole, 3-[(3,4-dichlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-,  
 monohydrochloride (9CI) (CA INDEX NAME)



● HCl

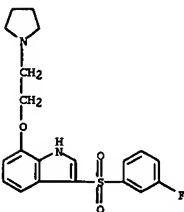
RN 706783-70-2 CAPLUS  
 CN 1H-Indole, 3-[(2-fluorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-,  
 monohydrochloride (9CI) (CA INDEX NAME)

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)



● HCl

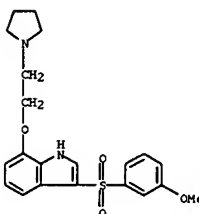
RN 706783-71-3 CAPLUS  
 CN 1H-Indole, 3-[(3-fluorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-,  
 monohydrochloride (9CI) (CA INDEX NAME)



● HCl

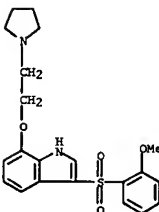
RN 706783-72-4 CAPLUS  
 CN 1H-Indole, 3-[(3-methoxyphenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-,  
 monohydrochloride (9CI) (CA INDEX NAME)

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)



● HCl

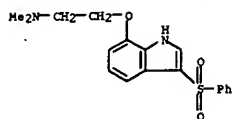
RN 706783-73-5 CAPLUS  
 CN 1H-Indole, 3-[(2-methoxyphenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-,  
 monohydrochloride (9CI) (CA INDEX NAME)



● HCl

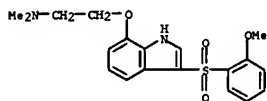
RN 706783-74-6 CAPLUS  
 CN Ethanamine, N,N-dimethyl-2-[[3-(phenylsulfonyl)-1H-indol-7-yl]oxy]-,  
 monohydrochloride (9CI) (CA INDEX NAME)

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)



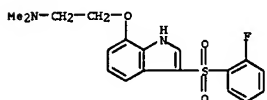
● HCl

RN 706783-75-7 CAPLUS  
 CN Ethanamine, 2-[[3-[(2-methoxyphenyl)sulfonyl]-1H-indol-7-yl]oxy]-N,N-dimethyl-, monohydrochloride (9CI) (CA INDEX NAME)



● HCl

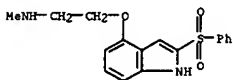
RN 706783-76-8 CAPLUS  
 CN Ethanamine, 2-[[3-[(2-fluorophenyl)sulfonyl]-1H-indol-7-yl]oxy]-N,N-dimethyl-, monohydrochloride (9CI) (CA INDEX NAME)



● HCl

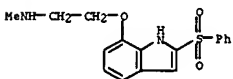
RN 706783-77-9 CAPLUS  
 CN Ethanamine, 2-[[3-[(2-fluorophenyl)sulfonyl]-1H-indol-7-yl]oxy]-N-methyl-, monohydrochloride (9CI) (CA INDEX NAME)

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)  
 CN Ethanamine, N-methyl-2-[[2-(phenylsulfonyl)-1H-indol-4-yl]oxy]-, monohydrochloride (9CI) (CA INDEX NAME)



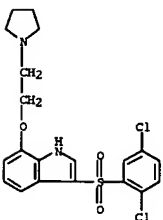
● HCl

RN 706783-83-7 CAPLUS  
 CN Ethanamine, N-methyl-2-[[2-(phenylsulfonyl)-1H-indol-7-yl]oxy]-, monohydrochloride (9CI) (CA INDEX NAME)



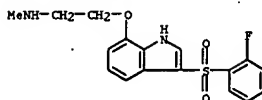
● HCl

RN 706783-84-8 CAPLUS  
 CN 1H-Indole, 3-[(2,5-dichlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-, monohydrochloride (9CI) (CA INDEX NAME)



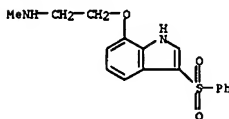
● HCl

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)



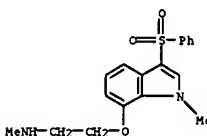
● HCl

RN 706783-78-0 CAPLUS  
 CN Ethanamine, N-methyl-2-[[3-(phenylsulfonyl)-1H-indol-7-yl]oxy]-, monohydrochloride (9CI) (CA INDEX NAME)



● HCl

RN 706783-79-1 CAPLUS  
 CN Ethanamine, N-methyl-2-[[1-methyl-3-(phenylsulfonyl)-1H-indol-7-yl]oxy]-, monohydrochloride (9CI) (CA INDEX NAME)

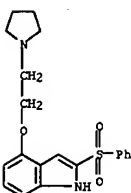


● HCl

RN 706783-82-6 CAPLUS

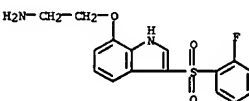
L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)

RN 706783-85-9 CAPLUS  
 CN 1H-Indole, 2-(phenylsulfonyl)-4-[2-(1-pyrrolidinyl)ethoxy]-, monohydrochloride (9CI) (CA INDEX NAME)



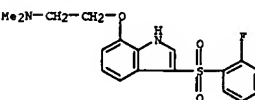
● HCl

RN 706783-87-1 CAPLUS  
 CN Ethanamine, 2-[[3-[(2-fluorophenyl)sulfonyl]-1H-indol-7-yl]oxy]-, monohydrochloride (9CI) (CA INDEX NAME)



● HCl

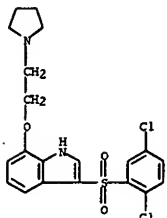
RN 706784-06-7 CAPLUS  
 CN Ethanamine, 2-[[3-[(2-fluorophenyl)sulfonyl]-1H-indol-7-yl]oxy]-N,N-dimethyl- (9CI) (CA INDEX NAME)



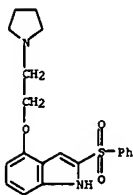
RN 706784-07-8 CAPLUS



L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)  
 CN 1H-Indole, 3-[(2,5-dichlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-  
 (9CI) (CA INDEX NAME)

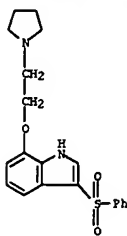


RN 706784-23-8 CAPLUS  
 CN 1H-Indole, 2-(phenylsulfonyl)-4-[2-(1-pyrrolidinyl)ethoxy]- (9CI) (CA INDEX NAME)

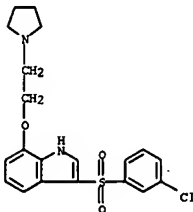


RN 706784-35-2 CAPLUS  
 CN 1H-Indole, 3-(phenylsulfonyl)-7-[2-(1-pyrrolidinyl)ethoxy]- (9CI) (CA INDEX NAME)

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)

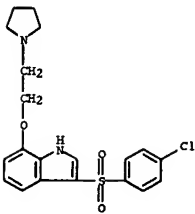


RN 706784-36-3 CAPLUS  
 CN 1H-Indole, 3-[(3-chlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-  
 (9CI) (CA INDEX NAME)

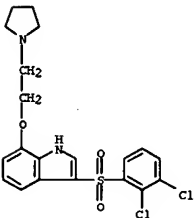


RN 706784-37-4 CAPLUS  
 CN 1H-Indole, 3-[(4-chlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-  
 (9CI) (CA INDEX NAME)

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)

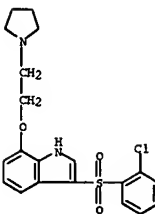


RN 706784-38-5 CAPLUS  
 CN 1H-Indole, 3-[(2,3-dichlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-  
 (9CI) (CA INDEX NAME)

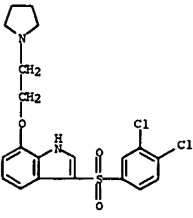


RN 706784-39-6 CAPLUS  
 CN 1H-Indole, 3-[(2-chlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-  
 (9CI) (CA INDEX NAME)

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)

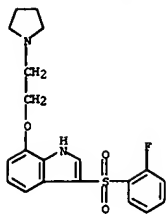


RN 706784-40-9 CAPLUS  
 CN 1H-Indole, 3-[(3,4-dichlorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-  
 (9CI) (CA INDEX NAME)

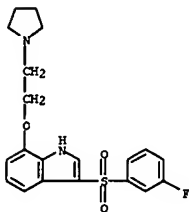


RN 706784-41-0 CAPLUS  
 CN 1H-Indole, 3-[(2-fluorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]-  
 (9CI) (CA INDEX NAME)

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)

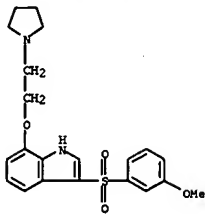


RN 706784-42-1 CAPLUS  
CN 1H-Indole, 3-[(3-fluorophenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]- (9CI) (CA INDEX NAME)

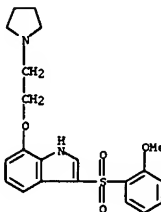


RN 706784-43-2 CAPLUS  
CN 1H-Indole, 3-[(2-methoxyphenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]- (9CI) (CA INDEX NAME)

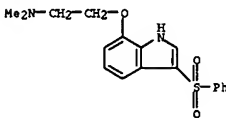
L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)



RN 706784-44-3 CAPLUS  
CN 1H-Indole, 3-[(2-methoxyphenyl)sulfonyl]-7-[2-(1-pyrrolidinyl)ethoxy]- (9CI) (CA INDEX NAME)

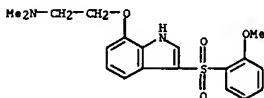


RN 706784-45-4 CAPLUS  
CN Ethanamine, N,N-dimethyl-2-[[3-(phenylsulfonyl)-1H-indol-7-yl]oxy]- (9CI) (CA INDEX NAME)

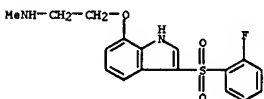


L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)

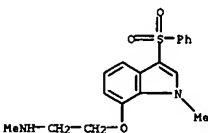
RN 706784-46-5 CAPLUS  
CN Ethanamine, 2-[[3-[(2-methoxyphenyl)sulfonyl]-1H-indol-7-yl]oxy]-N,N-dimethyl- (9CI) (CA INDEX NAME)



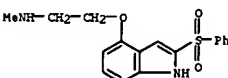
RN 706784-47-6 CAPLUS  
CN Ethanamine, 2-[[3-[(2-fluorophenyl)sulfonyl]-1H-indol-7-yl]oxy]-N-methyl- (9CI) (CA INDEX NAME)



RN 706784-48-7 CAPLUS  
CN Ethanamine, N-methyl-2-[[1-methyl-3-(phenylsulfonyl)-1H-indol-7-yl]oxy]- (9CI) (CA INDEX NAME)



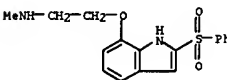
RN 706784-51-2 CAPLUS  
CN Ethanamine, N-methyl-2-[[2-(phenylsulfonyl)-1H-indol-4-yl]oxy]- (9CI) (CA INDEX NAME)



RN 706784-52-3 CAPLUS

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN (Continued)

RN 706784-49-8 CAPLUS  
CN Ethanamine, N-methyl-2-[[2-(phenylsulfonyl)-1H-indol-7-yl]oxy]- (9CI) (CA INDEX NAME)

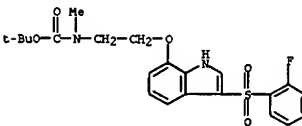


IT 706784-13-6P, [2-[[3-[(2-Fluorobenzesulfonyl)-1H-indol-7-yl]oxy]ethyl](methyl)carbamate] methyl ester 706784-17-0P, [2-[[3-[(2-Fluorobenzesulfonyl)-1-methyl-1H-indol-7-yl]oxy]ethyl](methyl)carbamate] methyl ester 706784-20-5P, 2-Benzenesulfonyl-4-[2-[(tert-butoxycarbonyl)(methyl)amino]ethoxy]indole-1-carboxylic acid tert-butyl ester 706784-22-7P, 2-Benzenesulfonyl-7-[2-[(tert-butoxycarbonyl)(methyl)amino]ethoxy]indole-1-carboxylic acid tert-butyl ester 706784-27-2P, 2-Benzenesulfonyl-4-[2-(pyrrolidin-1-yl)ethoxy]indole-1-carboxylic acid tert-butyl ester

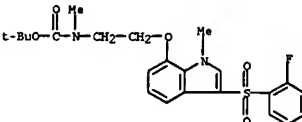
RL: RCT (Reactant); SPN (Synthetic preparation); PREF (Preparation); RACT (Reactant or reagent)

(intermediate; preparation of aminoalkoxyindoles as 5-HT6-receptor ligands, in particular selective 5-HT6 antagonists, for treating CNS disorders)

RN 706784-13-6 CAPLUS  
CN Carbamic acid, [2-[[3-[(2-fluorophenyl)sulfonyl]-1H-indol-7-yl]oxy]ethyl]methyl-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



RN 706784-17-0 CAPLUS  
CN Carbamic acid, [2-[[3-[(2-fluorophenyl)sulfonyl]-1-methyl-1H-indol-7-yl]oxy]ethyl]methyl-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



10724683 11/27/05

=> logoff

ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF

LOGOFF? (Y)/N/HOLD:y

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

10.33

171.87

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE

TOTAL

ENTRY

SESSION

CA SUBSCRIBER PRICE

-1.46

-1.46

STN INTERNATIONAL LOGOFF AT 14:07:58 ON 27 NOV 2005